

In the specification:

Please replace the paragraph at page 13, line 14 with the following:

PBMC were isolated from fresh whole blood by Ficoll density centrifugation. T-cells were purified by two rounds of treatment with Lympho-kwik (One Lambda, Canoga Park, CA). T-cell purity was verified by lack of a proliferative response to phytohemagglutinin ("PHA") or ~~{???~~ PMA in the absence of accessory cells. The human CD3-specific mAb HIT3a (PharMingen) was bound to 96-well plates at the indicated concentrations and used in this form to provide a first activating signal to T-cells. Alternatively, PHA was used in soluble form as a source of a first signal. K562 cells transfected with the negative control vector pREP7 $\beta$  (K562/pREP7 $\beta$ ) were precoated with pal-prot A and secondarily coated with B7-1-Fc $\gamma$ <sub>1</sub>. For each proliferation assay,  $1 \times 10^5$  T-cells were incubated with  $4 \times 10^4$  B7-1-Fc $\gamma$ <sub>1</sub>-coated and mitomycin C-treated K562/REP7 $\beta$  cells for 60 h at 37° C. Wells were pulsed with 1  $\mu$ Ci [<sup>3</sup>H]thymidine for the last 16 h of the incubation period. Cells were harvested and counted on a Betaplate liquid scintillation counter.